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Progress Report. Key Research Accomplishments, Reportable Outcomes, and Conclusions.

Introduction Update. PTEN has been characterized as a tumor suppressor gene and found to be deleted or mutated in a variety of human tumors, including breast cancer. PTEN functions with a dual-specificity protein and lipid phosphatase and negatively regulates cell growth, migration, invasion, and focal adhesions through down-regulation of downstream mediators, such as Akt, focal adhesion kinase, and Shr. Consistent with its tumor suppressor role, overexpression of PTEN in breast cancer cell lines may result in the suppression of the cell growth. Additionally, germline mutations in PTEN are associated with Cowden syndrome (CS), a rare autosomal disease characterized by an increased risk of developing breast cancer. On the other hand, androgen, acting through AR, has been suggested to inhibit mammary carcinoma growth in animal models and has been used clinically as an adjuvant therapeutic agent to influence breast cancer progression. In addition, the germline mutation of AR gene has been shown to cause partial androgen insensitivity together with familial male breast cancer, and the longer CAG repeat (>28) within AR gene (causing lower AR activity) combined with BRCA1 germline mutations is associated with an earlier age of onset of breast cancer development, suggesting that loss or reduction of AR function is implicated in the development of breast cancer. AR has been suggested to be importantly involved in breast cancer initiation. It also has been found that approximately 75%-80% of primary breast tumor specimens tested are positive for AR. Because PTEN expression is correlated with a positive ER and PR status in primary breast tumors and frequently reduced in advanced stage of breast cancer, it is conceivable to propose that PTEN counteracts AR in the mammary tissue to increase the susceptibility to transform to malignancy in the early stage of breast cancer development. Therefore, it will be interested to study the relationship between PTEN and AR in breast cancer and to determine if PTEN may modulate cell growth through AR.

Our studies have involved the MCF-7 breast cancer cell line and results are best summarized in the two figures presented below.

In Figure 1,we demonstrate that AR and PTEN are endogenously expressed in MCF7 cells. And we found that these two proteins can interact with each other in this cell line by co-immunoprecipitation.

As shown in Figure 2, using GST pull-down assay, we found that PTEN and AR are associated directly via the PTP domain of PTEN and AR DNA binding domain.

These studies were reported in a poster format at the Era of Hope Symposium in Orlando, Florida, September 25-28, 2002 and will be submitted along with other results to a peer reviewed Journal in the near future.

Figure 1. PTEN interacts with AR endogenously in the MCF-7 breast cancer cell line.

The MCF-7 cells were cultured in RPMI with 10% Charcoal-dextran treated fetal bovine serum media for 16 hours and treated with ethanol or 10 nM DHT for another 16 hours. Then, cells were harvested and subjected to co-immunoprecipitation with anti-PTEN and anti-HA (control) antibodies, followed by Western Blotting with AR and PTEN antibodies. Results shown are representative of several individual experiments.

Figure 2. GST or GST-PTEN incubation with the [35S]-labeled AR, ER, or RXR for 2 hours in the presence or absence of the ligand. The bound proteins were analyzed by SDS-PAGE, followed by autoradiography. B. Representation of PTEN deleted mutants. PTP domain, protein tyrosine phoshates domain; ty-p, tyrosine phosphorylation domain. C. [35S]-labeled AR was incubated with different PTEN deleted mutants. The nearly equivalent aliquots of PTEN deleted mutants used are shown in the right panel. D. Representation of AR deleted mutants. E. GST or GST-PTEN was incubated with different AR deleted mutants. The SDS-PAGE data represents results from several individual experiments.

Figure 1.

COIP:MCF-7

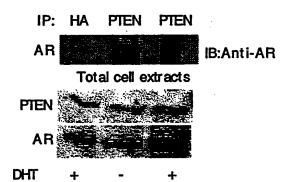


Figure 2.

PTEN interacts with AR in vitro.

